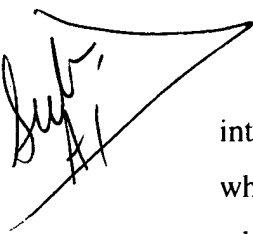
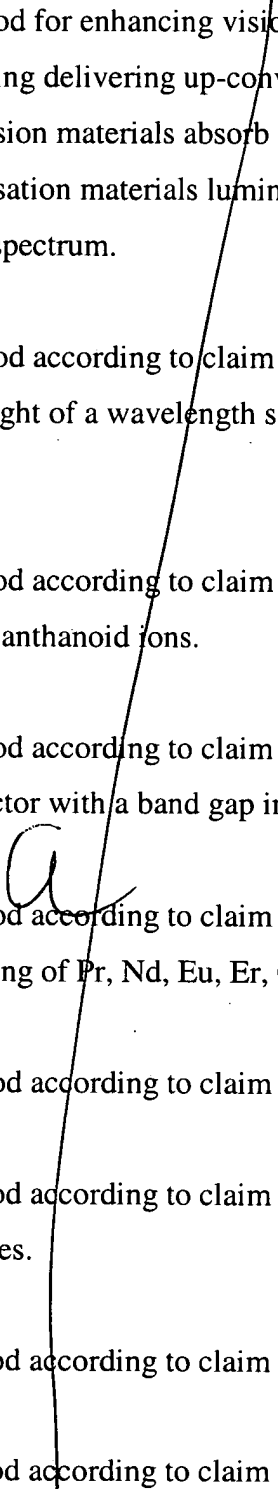


CLAIMS

I claim:

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1. A method for enhancing vision in an animal under conditions of low intensity light comprising delivering up-conversion materials to the eye of the animal, wherein the up-conversion materials absorb infrared light, and wherein the up-conversion materials luminescence in the visible range of the electromagnetic spectrum.
 2. A method according to claim 1, further comprising exposing the eye of the animal to a source of light of a wavelength sufficient to excite the up-conversion materials.
 3. A method according to claim 1, wherein the up-conversion materials comprise one or more lanthanoid ions.
 4. A method according to claim 1, wherein the up-conversion materials comprise a semiconductor with a band gap in the infrared.
 5. A method according to claim 3, wherein the lanthanoid ion is selected from the group consisting of Pr, Nd, Eu, Er, Gd, and Yb.
 6. A method according to claim 5, wherein the lanthanoid ion comprises Er.
 7. A method according to claim 1, wherein the up-conversion materials are in the form of nanoparticles.
 8. A method according to claim 7, wherein the nanoparticles comprise SiO₂.
 9. A method according to claim 7, wherein the nanoparticles comprise CdSe.

10. A method according to claim 1, wherein the up-conversion materials comprise a lanthanoid ion in a glass.
11. A method according to claim 7, wherein the nanoparticles are covalently bound to an antibody, wherein the antibody is specific for an antigen on a protein component of the eye.
12. A method according to claim 11, wherein the antibody is an antibody specific for a rod protein.
13. A method according to claim 11, wherein the antibody is specific for a cone protein.
14. A method according to claim 11, wherein the antibody is specific for ROM-1.
15. A method according to claim 11, wherein the antibody is specific for peripherin.
16. A method according to claim 11, wherein the antibody is specific for arrestin.
17. A method according to claim 11, wherein the antibody is specific for rhodopsin.
18. A method according to claim 1, wherein delivering the up-conversion material to the eye is carried out with iontophoresis.
19. A method according to claim 1, wherein the animal is a human.

20. A method according to claim 1, wherein the animal is non-human.
21. A composition comprising a nanoparticle covalently bound to an antibody, wherein the nanoparticle comprises an up-conversion material that absorbs electromagnetic radiation having a wavelength greater than about 650 nm and luminesces in the visible region of the electromagnetic spectrum, and the antibody is an antibody specific to a protein component of the eye.
22. A composition according to claim 21, wherein the antibody is specific to an antigen selected from the group consisting of rod proteins, cone proteins, ROM-1, peripherin, arrestin, S-antigen, and rhodopsin.
23. A composition according to claim 21, wherein the up-conversion material comprises one or more lanthanoid ions.
24. A composition according to claim 21, wherein the up-conversion material comprises a semiconductor having a band gap in the infrared.
25. A composition according to claim 21, wherein the nanoparticles comprise SiO₂.
26. A composition according to claim 21, wherein the nanoparticles comprise an organic polymer.
27. A composition according to claim 21, wherein the antibody is an antibody specific to peripherin.
28. A composition according to claim 21, wherein the antibody is an antibody specific to ROM-1.

29. An animal having enhanced vision, wherein an up-conversion material is optically coupled to the photoreceptors of at least one eye of the animal.

30. An animal according to claim 29, wherein the up-conversion of the material comprises nanoparticles comprising a material that absorbs infrared and luminesces visible light.

31. An animal according to claim 29, wherein the up-conversion material comprises one or more lanthanoid ions.

32. An animal according to claim 29, wherein the up-conversion material comprises two or more different lanthanoid ions.

33. An animal according to claim 29, wherein the up-conversion material comprises a semiconductor material having a band gap in the infrared.

34. An animal according to claim 29, wherein the up-conversion material is bound to an antibody that preferentially binds to a portion of one of the biomaterials in the eye.

35. An animal according to claim 34, wherein the antibody is an antibody to a rod protein.

36. An animal according to claim 34, wherein the antibody is an antibody to a cone protein.

37. An animal according to claim 34, wherein the antibody is an antibody to ROM-1.

38. An animal according to claim 34, wherein the antibody is an antibody to peripherin.

39. An animal according to claim 34, wherein the antibody is an antibody to X-arrestin.
40. An animal according to claim 34, wherein the antibody is an antibody to S-antigen.
41. An animal according to claim 34, wherein the antibody is an antibody to rhodopsin.
42. An animal according to claim 29, wherein the up-conversion material is optically coupled to two eyes of the animal.
43. A dog according to claim 29.
44. A method for visualizing an object under conditions of low ambient light comprising:
 exposing the object to incident electromagnetic radiation having a wavelength greater than what can be seen by the naked eye; and
 perceiving light reflected from the object with an enhanced eye,
wherein the enhanced eye comprises an up-conversion material optically coupled to the photoreceptors of the eye,
wherein the up-conversion material absorbs light of the wavelength reflected from the object, and luminesces in the visible region of the electromagnetic spectrum.
45. A method according to claim 44, wherein the up-conversion material comprises one or more lanthanoid ions.
46. A method according to claim 44, wherein the up-conversion material comprises two or more different lanthanoid ions.

47. A method according to claim 44, wherein the up-conversion material comprises a semiconductor having a band gap in the infrared.
48. A method according to claim 44, wherein the up-conversion material is in the form of a nanoparticle covalently bound to an antibody, wherein the antibody is specific for an antigen in a biomaterial found in the eye.
49. A method according to claim 48, wherein the antibody is an antibody to a rod protein.
50. A method according to claim 48, wherein the antibody is an antibody to a cone protein.
51. A method according to claim 48, wherein the antibody is an antibody to ROM-1.
52. A method according to claim 48, wherein the antibody is an antibody to peripherin.
53. A method according to claim 48, wherein the antibody is an antibody to S-antigen.
54. A method according to claim 48, wherein the antibody is an antibody to X-arrestin.
55. A method according to claim 44, wherein the incident electromagnetic radiation is light of a single frequency.
56. A method according to claim 44, wherein the incident electromagnetic radiation is coherent laser light.

57. A method according to claim 55, wherein the source of the light is a light emitting diode.

58. A method according to claim 44, wherein the object is continuously illuminated.

59. A method according to claim 44, wherein the object is illuminated by a source of non-classical light.

60. A method according to claim 44, further comprising providing a source of photons separate from the light reflected from the object, wherein the photons excite the up-conversion materials.

61. A method for visualizing an object with an enhanced eye, wherein the enhanced eye comprises an up-conversion material optically coupled to the photoreceptors of the eye, comprising
providing the eye with a first source of photons that sensitize the up-conversion material; and
providing the eye with a second source of photons reflected from the object, wherein the up-conversion material absorbs the light reflected from the object and luminesces in the visible.

62. A method according to claim 61, wherein the first source of photons is delivered to the eye without reflecting off the object.

63. A method according to claim 61, wherein the first source of photons has a wavelength of 1000 nm or less.

64. A method according to claim 61, wherein the second source of photons has a wavelength of 1500 nm or greater.

65. A method according to claim 61, wherein the second source of photons is from a CO₂ laser.

66. A method according to claim 61, wherein the first source of photons is provided by a light emitting diode.

67. A method according to claim 61, wherein the up-conversion material is in the form of nanoparticles.

68. A method according to claim 67, wherein the nanoparticle is covalently bound to an antibody for a protein component of the eye.

69. A method according to claim 67, wherein the antibody is an antibody specific for ROM-1 or peripherin.